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m/035/015

## Kennecott

July 2, 2001

Mr. Don A. Ostler Department of Environmental Quality Division of Water Quality P.O. Box 144870 Salt Lake City, Utah 84114-4870



Subject:

Net Acid Generation (NAG) Test Results for Humidity Cell Samples, Tailings Impoundment Ground Water Discharge Permit No. UGW350011

Dear Mr. Ostler:

In accordance with the compliance schedule of the Tailings Impoundment Groundwater Discharge Permit Number UGW350011 Kennecott Utah Copper (KUC) is required to perform kinetic net acid generation (NAG) testing on samples from the six humidity cells that were run in 1999 and 2000. The NAG tests have been completed and this letter provides a brief description of the results.

As shown on Table 1, the six humidity cell tailings samples were collected in 1999 from the Copperton Concentrator, Magna Concentrator and from the underflow stream at the cyclone stations. Neutralization potentials were relatively high in 1999 and only one of the six samples had a negative acid/base accounting potential. Sample HT9901 had an acid/base accounting potential of –17 and a neutralization potential ratio (NPR) of 0.4. All of the other samples had ABA potentials of 6 or higher and NPR values of 1.2 or higher. Whenever possible, the NAG tests were run on unmodified and untested tailings sample splits that had been archived, not on the material that had actually been placed in the humidity cell. However, for samples HT9901 and HT9904 no appropriate material was archived so the NAG tests had to be run on tailings that were taken from the humidity cells. The humidity cell test procedure partially oxidized the sulfides in this material and had removed some of the neutralization potential, so these two samples were reanalyzed by the modified Sobek acid/base accounting method as well as being NAG tested.

The temperature and pH results of the kinetic NAG tests are summarized on Table 1 and in Figures 1 through 6. Only the NAG test run on HT9901 acidified. The final pH of this

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test was 3.09 and the peak temperature was 65.98 degrees centigrade. It took almost 7 hours for the tailings to acidify. The leachate generated from HT9901 also had the highest sulfate, copper and zinc concentrations of any of the six tests (Table 2). The NAG test results for the sample are in good agreement with the ABA potential and NPR results which both predicted that HT9901 should acidify when exposed to surface weathering conditions. As shown on Table 3, the humidity cell for this sample never acidified despite being run for 47 weeks, about twice as long as typical tests.

None of the other five NAG tests acidified and all had final pH values above 7. The hydrogen peroxide solution typically has a pH of around 5.9 so in each test the pH dropped to around 6 when the solution was first mixed with the tailings sample. For every test except HT9901 the pH then gradually rose during the test. The temperature never exceeded 26 degrees C during any of these tests.

No tailings samples analyzed in a humidity cell by KUC have ever acidified. It is believed that the reaction kinetics in the humidity cells were very slow and that very few of the sulfides were ever oxidized. As expected, sulfide oxidation during the NAG tests is much more rapid. As shown on Table 2, between 27% and 100% of the sulfide sulfur in each sample was converted to sulfate and went into solution. These should be considered minimum values because a significant portion of the sulfate created by sulfide oxidation may have remained in the solid phase of the sample as secondary minerals.

All of the NAG tests were run for at least 24 hours to insure that the hydrogen peroxide was allowed to completely react with any available sulfides. The NAG test protocol states that the test should be run for a minimum of two hours and until all visible effervescing ceases. However sample HT9901 did not begin to acidify until several hours after visible effervescing had ended. To insure that future tests are not ended too soon, KUC recommends that the standard operating procedure (SOP) be modified to state that all tests will be run for 24 hours. It was also noted that when the leachate was being boiled at the end of the test, solids would sometimes precipitate out. In order to get representative concentrations of the dissolved constituents in the leachate, KUC recommends that the SOP text be changed to state that the liquid volume be maintained near 250 ml throughout the boiling process. After pH, conductivity, sulfate, alkalinity and acidity have been analyzed the leachate should then be digested to insure that all metals are returned to solution. KUC also suggests that the following analytical methods be cited in the revised SOP: pH – EPA 150.1, conductivity – EPA 120.1, acidity – Std. Method 2310B and Metals – EPA 200.8.

KUC intends to complete 15 more NAG tests during 2001. The samples selected for these tests will generally have an NPR of less than or equal to one, so it is anticipated that a much higher percentage of these future NAG tests will acidify.

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Should you have any questions or comments about these initial NAG test results, please call me at 569-7553 or Rich Borden at 569-7141.

Sincerely,

William J. Adams

Director, Environmental Affairs

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WJA:cl Enclosure

	- <u>.</u> 1						Paste	Initial	Final	Minutes	Initial	Highest	Final	Minutes to
Sample	Sampling	Sample		212 (4)	151 (0)	NDD (2)		рН	На	to pH <4	Temp.(5)	Temp.	Temp.	High Temp.
#	Location	Date	AP (1)	NP (1)		NPR (3)	pH (4)			405	22.2	65.9	23.4	585
HT9901(6)	Copperton	5/3/99	27	10	-17	0.4	7.19	6.02	3.09		21.3	25.9	24.6	388
HT9902	Magna	3/2/99	38	44	6	1.2	7.51	6.17	7.56	na			25.3	1036
HT9903	Underflow	6/30/99	38	59	21	1.6	8.22	6.12	7.53	na	21.5	26.3		942
HT9904 (6)	Copperton		7	35	28	2.4	6.93	6.37	7.33	na	21.7	25.2	24.3	
HT9905 (7)	Magna	9/13/99	19	26	7	1.4	8.31	6.00	7.14	na	21.9	25.8	23.8	811
		9/13/99	19	26	7	1.4	8.31	6.14	7.73	na	21.5	25.8	25.1	887
HT9906 (7)	Underliow	9/13/99	13			<del></del>			***					
	L			4°-1 /NIF	2)	eard in torn	os of tons	CaCO3	ner 1000	tons of taili	ngs.			
(1) Acid Pote	ntial (AP) an	d Neutraliz	ation Pot	ential (NF	are expre	ssea in tern	is or toris	Cacco	000	Tono or tain	ı.go.	<del>                                     </del>		
(2) The Acid/	(2) The Acid/Base Accounting (ABA) Potential equals NP minus AP.									<del> </del>		<del></del>		
(O) The News	(a) The Neutralization Detential Patio (NDP) equals NP divided by AP									<del>                                     </del>				
(4) The paste pH of the sample is measured by mixing distilled water with the tailings at a 1:1 ratio. The initial pH is measured after the hydrogen										<u> </u>				
peroxide (average pH = 5.9) has been added to the sample. The final pH is the pH at the end of the 24 hour test.														
(E) All tampo	ratura data i	c in dearee	e C		1	1				1	<u> </u>			
(5) All temperature data is in degrees C.  (6) No untested tailings sample was available for NAG testing HT9901 and HT9904. Tailings material that had already been partially oxidized in the										<u></u>				
(6) No untes	teu tailings s	ample was	available	101 11/10	l testing in it	1	T			1				
humidity cells	s had to be u	ised in thes	e NAG te	ests.		<u> </u>	1			0.000.000	gos except	for the NP	for samp	le .
(7) The ABA analyses for these samples were rerun. All AP, NP, ABA and NPR values for these samples are averages except for the NP for sample														
HT9906. Fo	HT9906. For this sample only the second NP result is reported because the first is believed to greatly underestimate NP.													

Sample			Total	Sulfide	Solution	% total S in	% Sulfide S	Solution	Solution	Solution
#	AP (1)	NP (1)	Sulfur (2)	Sulfur	Sulfate (3)	Solution (4)	in Solution (5)	Cu (3)	As	Zn
HT9901	27	10	0.87	0.87	118	100	100	12100	<5	244
HT9902	38	44	1.25	1.21	56	45	43	<20	84	<10
HT9903	38	59	1.30	1.27	37	28	27	36	62	34
HT9904	7	35	0.23	0.23	17	74	74	27	42	31 .
HT9905	19	26	0.71	0.65	39	55	51	<20	111	18
HT9906	19	26	0.74	0.63	54	73	68	28	59	48
(1) Acid Potential (AP) and Neutralization Potential (NP) are expressed in terms of tons CaCO3 per 1000 tons of tailings.										
(2) The total sulfur and the sulfide sulfur in the solid sample expressed in percent.										
(3) Sulfate in	(3) Sulfate in mg/L and metals in ug/L.									
(4) Based up	(4) Based upon the sulfate in solution and the mixing ratio of 250 grams of hydrogen peroxide solution for every 2.5 grams of tailings,									
percentage of the total sulfur removed from the sample can be calculated.										
(5) Assuming that all sulfate sulfur is removed from the sample first, this is the amount of sulfide sulfur that must have been										
converted to sulfate and then went into solution.										

Table 2 - Final NAG Leachate Dissolved Constituents and Estimates of Sulfur Removed from the Tailings Samples

Sample			Paste	Final Humidity	Final NAG	Final HC	NAG	Final HC	NAG
#	ABA (1)	NPR (2)	pH (3)	Cell (HC) pH	рΗ	Sulfate (4)	Sulfate	Alkalinity	Alkalinity
HT9901(5)	-17	0.4	7.90 (7.19)	6.50	3.09	50	118	5	-29
HT9902	6	1.2	7.51	6.94	7.56	55	56	12	111
HT9903	21	1.6	8.22	7.53	7.53	20	37	20	61
HT9904 (5)	28	2.4	7.67 (6.93)	7.34	7.33	60	17	15	100
HT9905	7	1.4	8.31	7.03	7.14	30	39	10	41
HT9906	7	1.4	8.31	7.52	7.73	27	54	19	81
			-						
(1) The Acid/Base Accounting (ABA) Potential equals NP minus AP.									
(2) The Neutralization Potential Ratio (NPR) equals NP divided by AP.									
(3) The paste pH of the sample is measured by mixing distilled water with the tailings at a 1:1 ratio. The humidity cell pH is the pH									
of the water dr	aining from	the cell durin	g the last we	ek of the test. Th	e final NAG pl	H is the pH of	the hydroger	n peroxide sol	ution at
the end of the	NAG test.								
(4) All sulfate concentrations in mg/L. All alkalinity values are in mg/L as CaCO3. A negative value indicates that the solution is									
acidic and is a	measure of	the acidity.							
(5) No fresh tailings sample was available for NAG testing HT9901 and HT9904. Tailings material that had already been partially									
oxidized in the humidity cells had to be used in these NAG tests. The first paste pH listed is for the sample before the humidity cell									
was run. The pH shown in parentheses was measured on the tailings after the humidity cell was completed and before the NAG									
test was initiate	ed.	-							

Table 3 - Comparison of Humidity Cell and NAG Test Data























